Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claims 1-32 are canceled.

- 36 33. (Currently Amended) A method of providing release of for treating obesity by releasing cholecystokinin peptide in a subject, comprising

 (A) administering to the subject an effective amount of a luminal cholecystokinin releasing factor polypeptide-oligomer conjugate, said conjugate comprising
 - i) a lysine residue;
 - ii) an oligomeric moiety attached to the N-terminus of the luminal cholecystokinin releasing factor polypeptide-oligomer conjugate; and
 - whereby upon administration to the subject, said <u>polypeptide-oligomer conjugate</u> compound integrates into a cell membrane of the gut epithelium of the subject wherein the luminal cholecystokinin releasing factor polypeptide-<u>oligomer conjugate</u> binds with a target receptor on the surface of an epithelial cell, thereby providing release of cholecystokinin <u>peptide</u>, and

(B) inducing satiety, whereby food intake is reduced.

- 37 34. (Currently Amended) The method of claim 33 36, wherein the oligomeric moiety attached to the N-terminus of the luminal cholecystokinin releasing factor peptide is a branched oligomeric moiety.
- 38 35. (Currently Amended) The method of claim 34 37, wherein the branched oligomeric moiety has the following formula:

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$$\begin{array}{c|c} Me(OCH_2CH_2)_nOCH_2(CH_2)_mCHCHNH & \\ & \\ & \\ Me(OCH_2CH_2)_nO \end{array}$$

where n is from 3 to 230 and m is from 0 to 20.

39 36. (Currently Amended) The method of claim 34 37, wherein the branched oligomeric moiety has the following formula:

where n is from 3 to 230 and m is from 0 to 20 and X is selected from the group consisting of N, O or S.

- 40 37. (Currently Amended) The method of claim 34 37, wherein the branched oligomeric moiety has a total average molecular weight of 4,000 to 10,000 Daltons.
- 41 38. (Currently Amended) The method of claim 33 36, wherein the oligomeric moiety is attached to the N-terminus using a hydrolyzable linker.
- 42 39. (Currently Amended) The method of claim 34 37, wherein the branched oligomeric moiety is attached to the N-terminus using a non-hydrolyzable linker.
- 43 –40. (Currently Amended) The method of claim 33 36, wherein the oligomeric moiety attached to the N-terminus of the luminal cholecystokinin releasing factor polypeptide has a total average molecular weight of 4,000 to 10,000 Daltons.

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- 44 –41. (Currently Amended) The method of claim 33 36, wherein the oligomeric moiety is attached to the lysine reside using a hydrolyzable bond.
- 45 -42. (Currently Amended) The method of claim 33 36, wherein the oligomeric moiety attached to the lysine reside is a linear oligomeric moiety.
- 46 -43. (Currently Amended) The method of claim -42 45, wherein the linear oligomeric moiety is attached to the lysine reside using a hydrolyzable bond.
- 47 -46. (Currently Amended) The method of claim 33 36, further comprising a lysine reside at the C-terminus of the luminal cholecystokinin releasing factor polypeptide.
- 48 -47. (Currently Amended) The method of claim -46 <u>47</u>, further comprising a linear oligomeric moiety attached to the lysine reside at the C-terminus of the luminal cholecystokinin releasing factor polypeptide.
 - 49 48. (Withdrawn)
 - 50 49. (Withdrawn)
 - 51 50. (Withdrawn)
 - 52 51. (Withdrawn)
 - 53 52. (Withdrawn)
 - 54 53. (Withdrawn)
 - 55 54. (Withdrawn)

- 56 55. (Withdrawn)
- 57 56. (Withdrawn)
- 58 57. (Withdrawn)
- 59 58. (Withdrawn)
- 60 59. (Withdrawn)
- 61 60. (Withdrawn)
- 62 61. (Currently Amended) A method of providing release of for treating obesity by releasing cholecystokinin peptide in a subject, comprising
- (A) administering to the subject an effective amount of a luminal cholecystokinin releasing factor polypeptide-oligomer conjugate, said conjugate comprising
 - i) a first lysine residue;
 - ii) a second lysine residue at the C-terminus of the luminal cholecystokinin releasing factor polypeptide-oligomer conjugate;
 - iii) a branched oligomeric moiety attached to the N-terminus of the luminal cholecystokinin releasing factor polypeptide-oligomer conjugate using an non-hydrolyzable linker;
 - iv) a linear oligomeric moiety attached to the first lysine reside of the luminal cholecystokinin releasing factor polypeptide-oligomer conjugate using a hydrolyzable bond; and
 - v) a linear oligomeric moiety attached to the second lysine reside at the Cterminus of the luminal cholecystokinin releasing factor polypeptideoligomer conjugate,

whereby, upon administration to the subject, said <u>polypeptide-oligomer</u>

<u>conjugate</u> compound integrates into a cell membrane of the gut epithelium of
the subject wherein the luminal cholecystokinin releasing factor polypeptide-<u>oligomer</u>

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<u>conjugate</u> binds with a target receptor on the epithelial cell surface, thereby providing release of cholecystokinin <u>peptide</u>, and

(B) inducing satiety, whereby food intake is reduced.

63 62. (Currently Amended) The method of claim 61 62, wherein the branched oligomeric moiety has the following formula:

$$Me(OCH_2CH_2)_nOCH_2(CH_2)_mCHCHNH$$
 $Me(OCH_2CH_2)_nO$

where n is from 3 to 230 and m is from 0 to 20.

64 63. (Currently Amended) The method of claim 61 62, wherein the branched oligomeric moiety has the following formula:

$$\begin{array}{c|c} \text{Me}(\text{OCH}_2\text{CH}_2)_n\text{XCH}_2(\text{CH}_2)_m\text{CHCHNH} \\ \hline \\ \text{Me}(\text{OCH}_2\text{CH}_2)_n\text{X} \end{array}$$

where n is from 3 to 230 and m is from 0 to 20 and X is selected from the group consisting of N, O or S.

65 64. (Currently Amended) The method of claim 61 62, wherein the branched oligomeric moiety has a total average molecular weight of 4,000 to 10,000 Daltons.

66 65. (Withdrawn)

67 66. (Withdrawn)

68 67. (Withdrawn)

69 68. (Withdrawn)

70 69. (Withdrawn)

- 71. (New) The method of claim 36, wherein administering to the subject comprises orally administering to the subject.
- 72. (New) The method of claim 62, wherein administering to the subject comprises orally administering to the subject.